AMENDMENTS TO THE CLAIMS

The following listing of the claims will replace all prior versions, and listings, of claims in the application.

What is claimed is:

- (currently amended) A method of inhibiting Vif multimerization in a subject by comprising administering an effective amount of a Vif antagonist to the subject, wherein the Vif antagonist binds to the multimerization domain within a Vif protein, and the Vif antagonist is a peptide comprising a PXP motif.
- 2. (withdrawn; currently amended). The method of claim 1, wherein the Vif antagonist is an anti-Vif antibody; or a Vif protein fragment which comprises at least SEQ ID NO: 25; or a peptide comprising a PXP motif.
- 3. (withdrawn) The method of claim 2, wherein the Vif antagonist is an anti-Vif antibody.
- 4. (withdrawn) The method of claim 2, wherein the Vif protein fragment is selected from the group consisting of SEQ ID NO: 1, SEQ ID NO: 21, SEQ ID NO: 22, SEQ ID NO: 23, and SEQ ID NO: 26.
- 5. (withdrawn) The method of claim 1, wherein the Vif antagonist is an analog or derivative of a Vif protein fragment which comprises at least SEQ ID NO: 25.
- 6 (withdrawn) The method of claim 1, wherein the Vif antagonist is a chimeric protein comprising a Vif protein, or a chimeric protein comprising a Vif protein fragment which comprises at least SEQ ID NO: 25.
- 7. (canceled).
- 8. (currently amended). The method of claim [7] 1, wherein X in the PXP motif is any one amino acid.

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- 9. (currently amended). The method of claim 8, wherein the any one amino acid <u>is</u> selected from the group consisting of Arg, Val, Pro, Ser, Leu, Phe, Ala, His and Tyr.
- 10. (withdrawn) The method of claim 7, wherein X in the PXP motif is any two amino acids.
- 11. (withdrawn) The method of claim 10, wherein the any two amino acids are Pro-Pro or Pro-Leu.
- 12. (currently amended) The method of claim [7]1, wherein the peptide[[s]] comprising the PXP motif is are 4 to 20 amino acids long.
- 13. (currently amended) The method of claim 12, wherein peptide[[s]] comprising the PXP motif is are 5 to 20 amino acids long.
- 14. (currently amended) The method of claim 12, wherein peptide[[s]] comprising the PXP motif is are 12 amino acids long.
- 15 (currently amended). The method of claim [[7]] 1, wherein the peptide[[s]] comprising the PXP motif is are selected from the group consisting of SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID NO: 7, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO: 10, SEQ ID NO: 11, SEQ ID NO: 12, SEQ ID NO: 13, SEQ ID NO: 14, SEQ ID NO: 15, SEQ ID NO: 16, SEQ ID NO: 17, SEQ ID NO: 18, SEQ ID NO: 19, and SEQ ID NO: 20.
- 16 (withdrawn) The method of claim 1, wherein the Vif antagonist is a peptidomimetic of a peptide comprising a PXP motif.
- 17. (original). The method of claim 1, wherein the Vif antagonist is a molecule comprising one or more peptides comprising a PXP motif.
- 18. (withdrawn) The method of claim 1, wherein the Vif antagonist is a molecule comprising one or more peptidomimetics of peptides comprising a PXP motif.

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- 19 (original) The method of claim 1, wherein the Vif antagonist is administered parenterally or enterally.
- 20 (original) The method of claim 1, wherein the Vif antagonist is administered locally.
- 21. (new) The method of claim 1, wherein the peptide comprising the PXP motif comprises the sequence SEQ ID NO: 11.
- 22. (new) The method of claim 21 wherein the peptide comprising the PXP motif consists of the sequence SEQ ID NO: 11.